=> fil reg

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## http://www.cas.org/ONLINE/UG/regprops.html

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1.1

L1 STR

L2 11 SEA SSS FUL L1

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FILE 'CAPLUS' ENTERED AT 11:24:49 ON 08 JAN 2007

L3 2 SEA ABB=ON PLU=ON L2

D SCAN TI

=> d que stat 12

L1 STR

Page 1-A

Page 2-A

J G1 8

Page 3-A VAR G1=X/H VAR G2=33/CY NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 26

STEREO ATTRIBUTES: NONE

L2 11 SEA FILE=REGISTRY SSS FUL L1

100.0% PROCESSED 16 ITERATIONS SEARCH TIME: 00.00.01

11 ANSWERS

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http://www.cas.org/ONLINE/UG/regprops.html

=> d que stat 12 L1 STR

Ak @33

12 N 10

Page 1-A

Page 2-A

61.8

Page 3-A VAR G1=X/H VAR G2=33/CY NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 26

STEREO ATTRIBUTES: NONE

L2 11 SEA FILE=REGISTRY SSS FUL L1

100.0% PROCESSED 16 ITERATIONS 11 ANSWERS

SEARCH TIME: 00.00.01

=> fil caplus FILE 'CAPLUS' ENTERED AT 11:26:00 ON 08 JAN 2007 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2007 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 8 Jan 2007 VOL 146 ISS 3 FILE LAST UPDATED: 7 Jan 2007 (20070107/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

http://www.cas.org/infopolicy.html 'OBI' IS DEFAULT SEARCH FIELD FOR 'CAPLUS' FILE

=> d que nos 13

L1STR

L2 11 SEA FILE=REGISTRY SSS FUL L1

L3 2 SEA FILE=CAPLUS ABB=ON PLU=ON L2

=> d .ca hitstr 13 1-2

ANSWER 1 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN 2004:654779 CAPLUS Full-text

ACCESSION NUMBER: DOCUMENT NUMBER:

141:190783

TITLE:

SOURCE:

Preparation of isoxazole-containing thiourea

inhibitors useful for treatment of varicella zoster

virus

INVENTOR(S):

Bloom, Jonathan David

PATENT ASSIGNEE(S):

Wyeth Holdings Corporation, USA U.S. Pat. Appl. Publ., 9 pp.

CODEN: USXXCO

DOCUMENT TYPE:

LANGUAGE:

Patent English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

]	PATENT NO.					KIND		DATE			APPLICATION NO.									
	US	S 2004157900									US 2004-772799									
		2004072052														20040209				
٧	WO	2004072052																		
		W:	ΑE,	ΑE,	AG,	AL,	AL,	ΑM,	AM,	AM,	ΑT,	ΑT,	ΑU,	ΑZ,	ΑZ,	BA,	BB,	BG,		
			BG,	BR,	BR,	BW,	BY,	BY,	BZ,	BZ,	CA,	CH,	CN,	CN,	CO,	CO,	CR,	CR,		
			CU,	CU,	CZ,	CZ,	DE,	DE,	DK,	DK,	DM,	DZ,	EC,	EC,	ΕĖ,	EE,	EG,	ES,		
			ES,	FI,	FI,	GB,	GD,	GE,	GE,	GH,	GM,	HR,	HR,	HU,	HU,	ID,	IL,	IN,		
								KG,												
								LU,												
			MZ,	MZ,	NA,	NI				•	•	•	•	•		•	•	•		
		RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AT,	BE,		
			BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,	IT,	LU,		
			MC,	NL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN.		
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				-	•	•	-	SN,		•	•	•	•	•	•	•	•	- •		
PRIOR	PRIORITY APPLN. INFO.:							US 2003-446602P								P 20030211				
OTHER	OTHER SOURCE(S):																			
	_		,				_													

ED Entered STN: 13 Aug 2004

GΙ

$$\mathbb{R}^{2} \mathbb{N}^{1}$$

The title compds. [I; R1 = halo, H; R2 = alkyl; X = II, III; R3 = alkyl, cycloalkyl, hydroxymethyl, etc.; R4 = alkyl which may be further substituted with (un)substituted Ph, cycloalkyl, pyridyl, etc.], useful for inhibiting replication of a herpes virus, were prepared E.g., a multi-step synthesis of 1-[4-(4-benzylisoxazol-3-yl)phenyl]-3-[1-(4-fluorophenyl)ethyl]thiourea (IV), was given. Seventeen title compds. I were prepared as described for IV, and tested for activity as herpes virus inhibitors (IC50 values against VZV, MTS, CMV, HSV and RSV were given).

IC ICM C07D261-02 ICS A61K031-42

INCL 514378000; 548240000

CC 28-6 (Heterocyclic Compounds (More Than One Hetero Atom))

Section cross-reference(s): 1, 63

IT 736982-12-0P 736982-13-1P 736982-14-2P 736982-15-3P 736982-16-4P

736982-17-5P 736982-18-6P 736982-19-7P

736982-20-0P 736982-21-1P 736982-22-2P

736982-23-3P 736982-24-4P 736982-25-5P

736982-26-6P 736982-27-7P 736982-28-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of isoxazole-containing thiourea inhibitors useful for treatment of

varicella zoster virus)

IT 736982-18-6P 736982-19-7P 736982-20-0P

736982-21-1P 736982-22-2P 736982-23-3P

736982-24-4P 736982-25-5P 736982-26-6P

736982-27-7P 736982-28-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of isoxazole-containing thiourea inhibitors useful for treatment of

varicella zoster virus)

RN 736982-18-6 CAPLUS

CN Thiourea, N-[1-(4-fluorophenyl)ethyl]-N'-[4-[3-(3-pyridinyl)-4-isoxazolyl]phenyl]- (9CI) (CA INDEX NAME)

RN 736982-19-7 CAPLUS

CN Thiourea, N-[1-(4-fluorophenyl)ethyl]-N'-[4-[3-(4-quinolinyl)-4-isoxazolyl]phenyl]- (9CI) (CA INDEX NAME)

RN 736982-20-0 CAPLUS

CN Thiourea, N-[1-(4-fluorophenyl)ethyl]-N'-[4-[3-(4-pyridinyl)-4-isoxazolyl]phenyl]- (9CI) (CA INDEX NAME)

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RN 736982-21-1 CAPLUS

CN Thiourea, N-[4-[3-(4-aminophenyl)-4-isoxazolyl]phenyl]-N'-[1-(4-fluorophenyl)ethyl]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

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RN 736982-22-2 CAPLUS

CN Thiourea, N-[1-(4-fluorophenyl)ethyl]-N'-[4-[3-(1,2,3-thiadiazol-4-yl)-4-isoxazolyl]phenyl]- (9CI) (CA INDEX NAME)

RN 736982-23-3 CAPLUS
CN Thiourea, N-[1-(4-fluorophenyl)ethyl]-N'-[4-[3-(2-pyridinyl)-4-isoxazolyl]phenyl]- (9CI) (CA INDEX NAME)

RN 736982-24-4 CAPLUS
CN Thiourea, N-[4-[3-[4-(dimethylamino)phenyl]-4-isoxazolyl]phenyl]-N'-[1-(4-fluorophenyl)ethyl]- (9CI) (CA INDEX NAME)

PAGE 2-A

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RN 736982-25-5 CAPLUS

CN Thiourea, N-[1-(4-fluorophenyl)ethyl]-N'-[4-[3-(4-hydroxyphenyl)-4-isoxazolyl]phenyl]- (9CI) (CA INDEX NAME)

#### PAGE 1-A

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RN 736982-26-6 CAPLUS

CN Thiourea, N-[1-(4-fluorophenyl)ethyl]-N'-[4-(3-phenyl-4-isoxazolyl)phenyl](9CI) (CA INDEX NAME)

RN 736982-27-7 CAPLUS

CN Thiourea, N-[1-(4-fluorophenyl)ethyl]-N'-[4-[3-(1H-imidazol-2-yl)-4-isoxazolyl]phenyl]- (9CI) (CA INDEX NAME)

PAGE 1-A

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RN 736982-28-8 CAPLUS

CN Thiourea, N-[1-(4-fluorophenyl)ethyl]-N'-[4-[3-(2-hydroxyphenyl)-4isoxazolyl]phenyl]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

PAGE 2-A

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ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 2004:581022 CAPLUS Full-text

DOCUMENT NUMBER: 141:292118

TITLE: Thiourea inhibitors of herpesviruses. Part 3:

Inhibitors of varicella zoster virus

Di Grandi, Martin J.; Curran, Kevin J.; Feigelson, AUTHOR(S):

Gregg; Prashad, Amar; Ross, Adma A.; Visalli, Robert; Fairhurst, Jeanette; Feld, Boris; Bloom, Jonathan D.

CORPORATE SOURCE: Wyeth Research, Pearl River, NY, 10965, USA

Bioorganic & Medicinal Chemistry Letters (2004), SOURCE:

14(16), 4157-4160

CODEN: BMCLE8; ISSN: 0960-894X

PUBLISHER: Elsevier Science B.V.

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 141:292118

ED Entered STN: 21 Jul 2004

AB The preparation of  $\alpha$ -methylbenzyl thioureas and their biol. activity against varicella zoster virus is described. Several analogs demonstrated IC50s < 0.1

 $\mu\text{M}$  and their SAR are discussed. These compds. represent a novel class of potent and selective nonnucleoside inhibitors of varicella zoster virus.

CC 10-5 (Microbial, Algal, and Fungal Biochemistry)

ΙT 62-56-6D, Thiourea, derivs. 273389-92-7 273390-57-1 273390-61-7 273392-13-5 273392-38-4 273392-57-7 273392-67-9 273392-78-2 273392-82-8 273394-77-7 273394-95-9 273394-97-1 273394-98-2 273395-10-1 273395-13-4 273395-24-7 273395-27-0 273395-29-2 273395-30-5 273395-31-6 273395-40-7 273395-37-2 273395-42-9 273395-55-4 273395-59-8 274262-04-3 274262-73-6 736982-12-0 736982-19-7 764723-07-1 764723-08-2 764723-09-3 764723-10-6 764723-11-7

RL: BSU (Biological study, unclassified); BIOL (Biological study) (thiourea inhibitors of varicella zoster virus)

IT 736982-19-7

RL: BSU (Biological study, unclassified); BIOL (Biological study) (thiourea inhibitors of varicella zoster virus)

RN 736982-19-7 CAPLUS

CN Thiourea, N-[1-(4-fluorophenyl)ethyl]-N'-[4-[3-(4-quinolinyl)-4-isoxazolyl]phenyl]- (9CI) (CA INDEX NAME)

REFERENCE COUNT:

=>

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